PATENT COOPERATION TREATY



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INTERNATIONAL PRELIMINARY EXAMINATION REPORT

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INTERNAT	IONAL PRELIMINAR	Y EXAMIN	ATION REPORT
	(PCT Article 36 as	nd.Rule 70)	
Applicant's or agent's file reference P 26826	FOR FURTHER ACTIO	See Notifi Preliminary	ication of Transmittal of Interna Examination Report (Form PCT/IPEA.
International application No. PCT/DE2003/003123	International filing date (day 19 September 2003 (1		Priority date (day/month/year) 19 September 2002 (19.09.20
International Patent Classification (IPC) or G01N 27/30	national classification and IPC		
Applicant	INFINEON TECHNO	LOGIES AG	}
This international preliminary example and is transmitted to the applicant.	mination report has been preparaccording to Article 36.	ed by this Inter	rnational Preliminary Examining Author
2. This REPORT consists of a total o	of 5 sheets, inclu	ding this cover	sheet.
amended and are the basis f	nied by ANNEXES, i.e., sheets for this report and/or sheets con ne Administrative Instructions u	taining rectific	tion, claims and/or drawings which have cations made before this Authority (see
These annexes consist of a	total of sheets		
3. This report contains indications re	lating to the following items:		
I Basis of the report	t		
II Priority			
III Non-establishmen	t of opinion with regard to nov	elty, inventive s	step and industrial applicability
IV Lack of unity of in	nvention		
V Reasoned stateme	nt under Article 35(2) with reg anations supporting such staten	ard to novelty, i nent	inventive step or industrial applicability
VI Certain document			
	the international application		
· · ·	ons on the international applica	tion	
Date of submission of the demand	Dat	e of completion	n of this report
19 April 2004 (19.04	4.2004)	25	January 2005 (25.01.2005)
Name and mailing address of the IPEA/E	P Au	horized officer	
Facsimile No.	Tel	ephone No.	

Form PCT/IPEA/409 (cover sheet) (July 1998)

International application No.

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PCT/DE2003/003123

I. Basis	of the rep	port					
1. With	regard to	the elements of the international application:*					
	the inter	national application as originally filed					
$\overline{\boxtimes}$	the desc	ription:					
	pages	1-27	, as originally filed				
	pages		, filed with the demand				
	pages	, filed with the letter of					
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	pages		, as originally filed				
	pages	, as amended (togethe	r with any statement under Article 19				
	pages		, filed with the demand				
i	pages	1-9, filed with the letter of	30 December 2004 (30.12.2004)				
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		wings. 1/4-4/4	, as originally filed				
1	pages	1/4-4/4	, filed with the demand				
•	pages	, filed with the letter of					
] [-	ence listing part of the description:	as originally filed				
l	pages		, filed with the demand				
	pages pages	, filed with the letter of					
3. Wife pred	 2. With regard to the language, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item. These elements were available or furnished to this Authority in the following language which is: the language of a translation furnished for the purposes of international search (under Rule 23.1(b)). the language of publication of the international application (under Rule 48.3(b)). the language of the translation furnished for the purposes of international preliminary examination (under Rule 55.2 and/or 55.3). 3. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing: contained in the international application in written form. filed together with the international application in computer readable form. furnished subsequently to this Authority in written form. furnished subsequently to this Authority in computer readable form. The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished. The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished. 						
in	This r	the description, pages the claims, Nos the drawings, sheets/fig report has been established as if (some of) the amendments had not been made, at the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).** at sheets which have been furnished to the receiving Office in response to an import as "originally filed" and are not annexed to this report since they do	vitation under Article 14 are referred to				
		ment sheet containing such amendments must be referred to under item 1 and a	nnexed to this report.				

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V.	Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability;
	citations and explanations supporting such statement

Statement			
Novelty (N)	Claims	1-9	YES
	Claims		NO NO
Inventive step (IS)	Claims	1-9	YES
	Claims		NO
Industrial applicability (IA)	Claims	1-9	YES
	Claims		NO

2. Citations and explanations

This report makes reference to the following document: D1: EP-A-0 299 778.

Claim 1:

D1 discloses a method for producing a biosensor circuit
arrangement (page 5)

- in which an integrated circuit is formed in a substrate (figures 1-7),
- in which a core of an integrated reference electrode is formed by printing on the substrate with silver material as metal (example 1),
- in which subsequently the core of silver material is surrounded, at least in part, by a sleeve made of a salt of the silver material that is not readily soluble, thereby forming the integrated reference electrode (example 1),
- in which the integrated circuit is electrically coupled with the core of the integrated reference electrode (trivial feature).

Therefore, the subject matter of claim 1 differs from the known method in that

- biological molecules are applied to sensor fields of

the biosensor circuit arrangement by means of printing, whereby the sensor fields are biologically activated and whereby the printing of silver material onto the substrate and the printing of biological molecules onto the sensor fields takes place in the same operational step.

In **D1**, the reference electrode is produced in its entirety before the biological molecules are applied in a further step (see example 2).

Therefore, the subject matter of claim 1 is novel (PCT Article 33(2)).

Consequently, the problem to be solved by the present invention can be regarded as that of significantly reducing the complexity and cost of producing biosensors, since the same printing method is used in a method step in order to apply the reference electrode as that used for the biomolecules. Thereafter, the silver core is oxidized in another method step.

Therefore, the solution to this problem as proposed in claim 1 of the present application involves an inventive step (PCT Article 33(3)).

Claim 2:

Like claim 1, claim 2 contains the additional feature

that biological molecules are applied to sensor fields of the biosensor circuit arrangement by means of printing, whereby the sensor fields are biologically activated and whereby the printing of silver material onto the substrate and the printing of biological molecules onto the sensor fields takes

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place in the same operational step.

Therefore, the solution to this problem as proposed in claim 2 of the present application involves an inventive step (PCT Article 33(3)).

Dependent claims 3-9 are therefore also regarded as novel and inventive.